Amendments To The Claims

1-23. (Canceled)

24. (Currently amended) A method of inducing B cell depletion in a patient in need of such depletion having a B cell disorder comprising administering a B cell depleting effective amount of a non-radiolabeled chimeric anti-CD20 antibody, wherein said chimeric anti-CD20 antibody wherein when administered by itself at a dosage of 0.4 mg/kg body weight results in merely complete B cell depletion of greater than 90% of peripheral B cells within about 24 hour hours post treatment infusion of said chimeric anti-CD20 antibody.

25-30. (Canceled)

- 31. (Original) The method of claim 24 wherein said chimeric anti-CD20 antibody contains the variable heavy sequence corresponding to SEQ ID NO: 11.
- 32. (Original) The method of claim 24 wherein said chimeric anti-CD20 antibody contains the variable light sequence corresponding to SEQ ID NO: 7.
- 33. (Original) The method of claim 24 which further includes the administration of at least one chemotherapeutic agent.
- 34. (Original) The method of claim 33 wherein said at least one chemotherapeutic agent is selected from the group consisting of cyclophosphamide, doxorubicin, vincristine and prednisone.

35-40. (Canceled)

41. (Original) The method of claim 24 which further includes the administration of a radiolabeled anti-CD20 antibody.

U.S. Appl. No. 08/921,060 Attorney Docket No. 037003-0275463

- 42. (Original) The method of claim 41 wherein said radiolabeled anti-CD20 antibody is a murine anti-CD20 antibody.
- 43. (New) The method of claim 24 wherein said chimeric anti-CD20 antibody contains the variable heavy sequence corresponding to SEQ ID NO: 11 and the variable region light sequence corresponding to SEQ ID NO:7, or CD20 binding fragment thereof.
- 44. (New) The method of claim 24 wherein said chimeric anti-CD20 antibody contains the complementarity determining regions of SEQ ID NOs: 7 and 11.